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## WHAT IS CLAIMED IS:

- 1. An isolated nucleic acid molecule comprising a nucleotide sequence selected from the group consisting of:
  - a) the nucleotide sequence as set forth in Figure 1A (SEQ ID NO: 1);
  - b) the nucleotide sequence encoding the polypeptide from residues 1-200 or from residues 21-200 as set forth in Figure 1A (SEQ ID NO: 1);
  - c) a nucleotide sequence encoding a polypeptide that is at least about 70 percent identical to the polypeptide as set forth in Figure 1A (SEQ ID NO: 1);
- d) a naturally occurring allelic variant or alternate splice variant of any of (a), (b) or (c);
  - e) a nucleotide sequence complementary to any of (a), (b) or (c);
  - f) a nucleotide sequence of (b),(c) or (d) encoding a polypeptide fragment of at least about 25, 50, 75, 100, or greater than 100 amino acid residues;
    - g) a nucleotide sequence of (a), (b) or (c) comprising a fragment of at least about 10, 15, 20, 25, 50, 75, 100, or greater than 100 nucleotides; and
- 25 h) a nucleotide sequence which hybridizes under stringent conditions to any of (a)-(g).
- 2. An isolated nucleic acid molecule comprising a nucleotide sequence selected from the group consisting 30 of:
  - a) the nucleotide sequence as set forth in Figure 2A (SEQ ID NO: 11) or Figure 3A (SEQ ID NO: 6) or Figure 12A (SEQ ID NO: 16);
- b) the nucleotide sequence encoding the 35 polypeptide as set forth in Figure 2A (SEQ ID NO: 6) from residues 1-322 or from residues 47-322, or as set

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forth in Figure 3A (SEQ ID NO: 11) from residues 1-288 or from residues 19-288, 20-288, 21-288, 22-288, 24-288, or 28-288 or as set forth in Figure 12A from residues 1-302, or from residues 19-302, 20-302, 21-302, 22-302, 24-302 or 28-302;

- c) a nucleotide sequence encoding a polypeptide that is at least about 70 percent identical to the polypeptide as set forth in Figure 2A (SEQ ID NO: 6) or Figure 3A (SEQ ID NO: 11) or Figure 12A (SEQ ID NO: 6);
- d) a naturally occurring allelic variant or alternate splice variant of any of (a), (b) or (c);
- e) a nucleotide sequence complementary to any
  of (a), (b) or (c);
- f) a nucleotide sequence of (b),(c) or (d) encoding a polypeptide fragment of at least about 25, 50, 75, 100, or greater than 100 amino acid residues;
  - g) a nucleotide sequence of (a), (b) or (c) comprising a fragment of at least about 10, 15, 20, 25,
- 20 50, 75, 100, or greater than 100 nucleotides; and

  h) a nucleotide sequence which hybridizes under stringent conditions to any of (a)-(g).
- The nucleic acid molecule of Claims 1 or 2
   wherein the nucleotide sequence is operably linked to an expression control sequence.
  - 4. A host cell comprising the nucleic acid molecule of Claim 2.
  - 5. The host cell of Claim 3 which is a eucaryotic cell.
- 6. The host cell of Claim 3 which is a procaryotic 35 cell.

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7. A process for producing a polypeptide comprising growing a culture of the host cell of Claim 3 in suitable culture medium and isolating the polypeptide from the culture.

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- 8. A polypeptide produced by the process of Claim 7.
- 9. A polypeptide encoded by the nucleic acid 10 molecule of Claim 1.
  - 10. A polypeptide encoded by the nucleic acid molecule of Claim 2.
- 15 11. An isolated polypeptide comprising the amino acid sequence selected from the group consisting of:
  - a) the amino acid sequence as set forth in Figure 1A (SEQ ID NO: 2);
- b) the mature amino acid sequence as set 20 forth in Figure 1A (SEQ ID NO: 2) comprising a mature amino terminus at residue 21;
  - c) a fragment of the amino acid sequence set forth in Figure 1A (SEQ ID NO: 2) comprising at least about 25, 50, 75, 100, or greater than 100 amino acid residues:
    - d) an ortholog of (a), (b) or (c); and
  - e) an allelic variant or alternative splice variant of (a), (b), (c) or (d).
- 30 12. An isolated polypeptide comprising the amino acid sequence selected from the group consisting of:
  - a) the amino acid sequence as set forth in Figure 2A (SEQ ID NO: 7) or Figure 3A (SEQ ID NO: 12) or Figure 12A (SEQ ID NO: 17);
- 35 b) the mature amino acid sequence as set forth in Figure 2A (SEQ ID NO: 7) comprising a mature

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amino terminus at residues 47, or Figure 3A (SEQ ID NO: 12) comprising a mature amino terminus at any of residues 19, 20, 21, 22, 24 or 28, or Figure 12A (SEQ ID NO: 17) comprising a mature amino terminus at any of residues 19,20,21,22,24,or 28;

- c) a fragment of the amino acid sequence set forth in Figure 2A (SEQ ID NO: 7) or Figure 3A (SEQ ID NO: 12) or Figure 12A (SEQ ID NO: 17) comprising at least about 25, 50, 75, 100, or greater than 100 amino acid residues;
- d) an ortholog of (a), (b) or (c); and
   e) an allelic variant or alternative splice
   variant of (a), (b), (c) or (d).
- 13. An antibody or fragment thereof specifically binding the polypeptide of Claims 9, 10, 11 or 12.
  - 14. The antibody of Claim 11 which is a monoclonal antibody.
  - 15. The antibody of Claim 13 which is a human antibody.
- 16. The antibody of Claim 13 which is a humanized or CDR-grafted antibody.
  - 17. The antibody or fragment of Claim 13 which binds B7RP1 or to a B7RP1 extracellular domain.
- 30 18. The antibody or fragment of Claim 13 which inhibits the binding of B7RP1 to CRP1.
- 19. A composition comprising the polypeptide of Claims 9, 10, 11 or 12 and a pharmaceutically35 acceptable carrier, adjuvant, solubilizer, stabilizer or anti-oxidant.

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- 20. A polypeptide comprising a derivative of the polypeptide of Claims 9, 10, 11 or 12.
- 5 21. The polypeptide of Claim 20 which is covalently modified with a water-soluble polymer.
  - 22. A fusion polypeptide comprising the polypeptide of Claims 9, 10, 11 or 12 fused to a heterologous amino acid sequence.
    - 23. The fusion polypeptide of Claim 22 wherein the heterologous amino acid sequence is an IgG constant domain or fragment thereof.
    - 24. A method for treating, preventing or ameliorating a T cell mediated disorder comprising administering to an animal the polypeptide of Claims 9, 10, 11 or 12.
    - 25. A method of diagnosing a T cell mediated disorder or a susceptibility to a T cell mediated disorder in an animal comprising:
- a) determining the presence or amount of expression of the polypeptide of Claims 9, 10, 11 or 12: and
  - b) diagnosing a T-cell mediated disorder or a susceptibility to a T-cell mediated disorder based on the presence or amount of expression of the polypeptide.
  - 26. A method of identifying a test molecule which binds to a polypeptide comprising:
- a) contacting the polypeptide of Claims 9,35 10, 11 or 12 with a test molecule; and

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- b) determining the extent of binding of the polypeptide to the test molecule.
- 27. The method of Claim 26 further comprising
  5 determining the activity of the polypeptide when bound to the compound.
- 28. A method of regulating T cell activation or proliferation in an animal comprising administering to the animal the nucleic acid molecule of Claims 1, 2 or 3.
  - 29. A transgenic non-human mammal comprising the nucleic acid molecule of Claim 3.
  - 30. A method of suppressing an immune response in an animal comprising administering to the animal an antagonist of CRP-1 or B7RP-1.
- 31. The method of Claim 30 wherein the antagonist is an antibody which binds B7RP-1.